



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C03C 3/087, 4/02, 4/08	A1	(11) International Publication Number: WO 99/20577 (43) International Publication Date: 29 April 1999 (29.04.99)
(21) International Application Number: PCT/US98/21987 (22) International Filing Date: 16 October 1998 (16.10.98) (30) Priority Data: 08/954,722 20 October 1997 (20.10.97) US 09/058,381 9 April 1998 (09.04.98) US (71) Applicant: PPG INDUSTRIES OHIO, INC. [US/US]; 3800 West 143rd Street, Cleveland, OH 44111 (US). (72) Inventor: SHELESTAK, Larry, J.; 1046 Ford Street, P.O. Box 233, Bairdford, PA 15006 (US). (74) Agents: STACHEL, Kenneth, J.; PPG Industries, Inc., One PPG Place, Pittsburgh, PA 15272 (US) et al.		(81) Designated States: AT, AU, BR, CA, CH, CN, DE, DK, ES, FI, GB, JP, KR, LU, MX, NZ, PT, SE, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i>
(54) Title: INFRARED AND ULTRAVIOLET RADIATION ABSORBING BLUE GLASS COMPOSITION (57) Abstract The present invention provides a blue colored glass using a standard soda-lime-silica glass base composition and additionally iron and cobalt, and optionally chromium, as solar radiation absorbing materials and colorants. In particular, the blue colored glass includes about 0.40 to 1.0 wt.% total iron, preferably about 0.50 to 0.75 wt.%, about 4 to 40 PPM CoO, preferably about 4 to 20 PPM, and 0 to 100 PPM Cr ₂ O ₃ . The redox ratio for the glass of the present invention is greater than 0.35 up to about 0.60, and preferably between about 0.36 to 0.50. In one particular embodiment of the invention, the glass has a luminous transmittance of at least 55 % and a color characterized by a dominant wavelength of 485 to 489 nanometers and an excitation purity of about 3 to 18 per cent. In another embodiment of the invention, the glass has a luminous transmittance of at least 65 % at a thickness of about 0.154 inches (3.9 mm) and a color characterized by a dominant wavelength of 485 to 492 nanometers and an excitation purity of about 3 to 18 percent.		

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INFRARED AND ULTRAVIOLET RADIATION ABSORBING BLUE GLASS COMPOSITION

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CROSS REFERENCE TO RELATED APPLICATION

This application is a continuation-in-part of U.S. Application Serial No. 08/954,722 filed October 20, 1997.

10

BACKGROUND OF THE INVENTION

1. Field of the Invention

This invention relates to an infrared and ultraviolet radiation absorbing blue colored soda-lime-silica glass composition suitable for architectural and automotive glazing applications. The glass should have a dominant
15 wavelength of about 485 to 492 nanometers (nm) and an excitation purity of about 3 to 18 percent. The glass is also compatible with flat glass manufacturing methods.

2A. Technical Considerations

Infrared and ultraviolet radiation absorbing colored glass substrates
20 have a variety of different applications. In particular, such glasses may be used by architects to glaze buildings and by vehicle designers as automotive windows. Besides providing an aesthetically pleasing color, these glasses may also provide enhanced solar performance as compared to conventional clear glass.

25 Different materials may be added to the glass in order to provide the desired color and spectral performance. For example, iron, cobalt, nickel, selenium and titanium, to name a few, are typically added to provide the desired color composition. As materials are added to change color and enhance solar performance, care must be taken to maintain the visible light
30 transmittance and color required for that particular application. It should also be remembered that changing the thickness of the glass will affect these spectral properties so that a particular composition which has acceptable

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color and performance at a particular thickness may not be acceptable at a different thickness.

One particular blue composition that provides superior spectral performance is disclosed in U.S. Patent No. 4,792,536 to Pecoraro, et al.

5 Commercial products which incorporate this patent are sold by PPG Industries, Inc. under the trademarks SOLEXTRA® and AZURLITE®. This glass incorporates a moderate amount of iron in the composition and has a relatively large portion of the glass in the ferrous state, expressed as FeO. In particular, the glass composition includes a basic soda-lime-silica composition
10 and further includes 0.45 to 1 weight percent total iron (expressed as Fe₂O₃). At least 35 percent of the total iron is in the ferrous state. The dominant wavelength of these glasses range from about 486 to 489 nm and excitation purity ranges from about 8 to 14 percent. From a processing standpoint, producing the glass disclosed in the patent with a high ratio of ferrous iron to
15 total iron may require additional processing considerations not typically associated with conventional glass melting operations, as are well known in the art. However, due to the favorable acceptance of this product, it would be advantageous to be able to produce a glass having a similar color and enhanced spectral performance using conventional glass melting processing
20 techniques.

2B. Patents of Interest

U.S. Patent No. 3,652,303 to Janakirama Rao discloses a blue, heat absorbing glass which incorporates low amounts of iron and uses tin to convert and retain a significant portion of the iron in the ferrous state, and in
25 particular more than 80% of the iron is retained in the ferrous state.

U.S. Patent Nos. 4,866,010 and 5,070,048 to Boulos, et al. disclose blue glass compositions with a colorant portion consisting essentially of iron and cobalt and further including nickel and/or selenium. The glasses have a

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dominant wavelength of 482 nanometers (nm) ± 1 nm and a color purity of 13% ± 1 %.

U.S. Patent Nos. 5,013,487 and 5,069,826 to Cheng disclose blue colored glass compositions which include iron, titanium, tin and zinc as
5 colorants. The glasses have a dominant wavelength ranging from 485 to 494 nm and a color purity of 5 to 9%.

U.S. Patent No. 5,344,798 to Morimoto, et al. discloses a blue glass composition which includes iron, cerium, titanium, zinc, cobalt and manganese. These glasses have a dominant wavelength of 495 to 505 nm
10 and a color purity of 5 to 9%.

SUMMARY OF THE INVENTION

The present invention provides a blue colored glass using a standard soda-lime-silica glass base composition and additionally iron and cobalt, and
15 optionally chromium, as solar radiation absorbing materials and colorants. In particular, the blue colored glass includes about 0.40 to 1.0 wt.% total iron, preferably about 0.50 to 0.75 wt.%, about 4 to 40 PPM CoO, preferably about 4 to 20 PPM, and 0 to 100 PPM Cr₂O₃. The redox ratio for the glass of the present invention is greater than 0.35 up to about 0.60, and preferably
20 between about 0.36 to 0.50. In one particular embodiment of the invention, the glass has a luminous transmittance of at least 55 percent and a color characterized by a dominant wavelength of 485 to 489 nanometers and an excitation purity of about 3 to 18 percent. In another embodiment of the invention, the glass has a luminous transmittance of at least 65 percent at a
25 thickness of about 0.154 inches (3.9 mm) and a color characterized by a dominant wavelength of 485 to 492 nanometers and an excitation purity of about 3 to 18 percent.

DETAILED DESCRIPTION OF THE INVENTION

The base glass composition of the present invention is commercial soda-lime-silica glass characterized as follows:

		<u>Weight Percent</u>
5	SiO ₂	66 to 75
	Na ₂ O	10 to 20
	CaO	5 to 15
	MgO	0 to 5
	Al ₂ O ₃	0 to 5
10	K ₂ O	0 to 5

As used herein, all "weight percent (wt.%) " values are based on the total weight of the final glass composition.

To this base glass, the present invention adds infrared and ultraviolet radiation absorbing materials and colorants in the form of iron and cobalt, and, optionally, chromium. As disclosed herein, iron is expressed in terms of Fe₂O₃ and FeO, cobalt is expressed in terms of CoO, and chromium is expressed in terms of Cr₂O₃. It should be appreciated that the glass compositions disclosed herein may include small amounts of other materials, for example melting and refining aids, tramp materials or impurities. It should be further appreciated that in one embodiment of the invention, small amounts of additional materials may be included to color the glass and/or improve its solar performance, as will be discussed later in more detail.

The iron oxides in a glass composition perform several functions. Ferric oxide, Fe₂O₃, is a good ultraviolet radiation absorber and operates as a yellow colorant in the glass. Ferrous oxide, FeO, is a good infrared radiation absorber and operates as a blue colorant. The total amount of iron present in the glasses disclosed herein is expressed in terms of Fe₂O₃ in accordance with standard analytical practice, but that does not imply that all of the iron is

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actually in the form of Fe_2O_3 . Likewise, the amount of iron in the ferrous state is reported as FeO , even though it may not actually be present in the glass as FeO . In order to reflect the relative amounts of ferrous and ferric iron in the glass compositions disclosed herein, the term "redox" shall mean the amount
5 of iron in the ferrous state (expressed as FeO) divided by the amount of total iron (expressed as Fe_2O_3). Furthermore, unless stated otherwise, the term "total iron" in this specification shall mean total iron expressed in terms of Fe_2O_3 , the term " Fe_2O_3 " shall mean iron in the ferric state expressed in terms of Fe_2O_3 and the term " FeO " shall mean iron in the ferrous state expressed in
10 terms of FeO .

CoO operates as a blue colorant and a weak infrared radiation absorber in the glass. Cr_2O_3 may be added to impart a green color component to the glass composition. In addition, it is believed that the chromium may also provide some ultraviolet radiation absorption. A proper
15 balance between the iron, i.e. ferric and ferrous oxides, and cobalt, and optionally chromium, content is required to obtain a glass with the desired blue color and spectral properties.

The glass of the present invention may be melted and refined in a continuous, large-scale, commercial glass melting operation and formed into
20 flat glass sheets of varying thickness by the float method in which the molten glass is supported on a pool of molten metal, usually tin, as it assumes a ribbon shape and is cooled in a manner well known in the art.

Although it is preferred that the glass disclosed herein be made using a conventional, overhead fired continuous melting operation, as is well known in
25 the art, the glass may also be produced using a multi-stage melting operation, as disclosed in U.S. Patent Nos. 4,381,934 to Kunkle, et al.; 4,792,536 to Pecoraro, et al. and 4,886,539 to Cerutti, et al. If required, a stirring arrangement may be employed within the melting and/or forming stages of

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the glass production operation to homogenize the glass in order to produce glass of the highest optical quality.

Depending on the type of melting operation, sulfur may be added to the batch materials of a soda-lime-silica glass as a melting and refining aid.

- 5 Commercially produced float glass may include up to about 0.3 wt.% SO_3 . In a glass composition that includes iron and sulfur, providing reducing conditions may create amber coloration which lowers luminous transmittance as discussed in U.S. Patent No. 4,792,536 to Pecoraro, et al. However, it is believed that the reducing conditions required to produce this coloration in
- 10 float glass compositions of the type disclosed herein are limited to approximately the first 20 microns of the lower glass surface contacting the molten tin during the float forming operation, and to a lesser extent, to the exposed upper glass surface. Because of the low sulfur content and the limited region of the glass in which any coloration could occur, depending on
- 15 the particular soda-lime-silica-glass composition, sulfur in these surfaces has no material effect on the glass color or spectral properties.

- It should be appreciated that as a result of forming the glass on molten tin as discussed above, measurable amounts of tin oxide may migrate into surface portions of the glass on the side contacting the molten tin. Typically,
- 20 a piece of float glass has an SnO_2 concentration of at least 0.05 to 2 wt.% in the first 25 microns below the surface of the glass that was in contact with the tin. Typical background levels of SnO_2 may be as high as 30 parts per million (PPM). It is believed that high tin concentrations in about the first 10 angstroms of the glass surface supported by the molten tin may slightly
- 25 increase the reflectivity of that glass surface; however, the overall impact on the glass properties is minimal.

Table 1 illustrates experimental glass melts which embody the principles of the present invention. Similarly, Table 2 illustrates a series of

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computer modeled glass compositions embodying the principles of the present invention. The modeled compositions were generated by a glass color and spectral performance computer model developed by PPG Industries, Inc. Table 3 illustrates the composition of several glass samples
5 incorporating the principles of the present invention, which were fabricated on a commercial, conventional, overhead fired continuous glass melter of the type disclosed earlier. Only the iron and cobalt portions of the compositions are listed in Tables 1 and 3 while Table 2 includes the iron, cobalt and chromium portions of the compositions. Tables 4-6 list the spectral properties
10 of the compositions shown in Tables 1, 2 and 3 at reference thicknesses of 0.084 inches (2.13 mm), 0.154 inches (3.9 mm) and 0.223 inches (5.66 mm), respectively.

With respect to the data provided in Tables 4-6 for Examples 1-16 of Table 1 and Examples 30-34 of Table 3, the luminous transmittance (LTA) is
15 measured using C.I.E. standard illuminant "A" with a C.I.E. 2° observer over the wavelength range of 380 to 770 nanometers. The solar transmittance (LTS) is measured over the wavelength range of 380 to 770 nm using the C.I.E. 2° observer and the weighting factors specified in ASTM 891-87. Glass color, in terms of dominant wavelength (DW) and excitation purity (Pe), is
20 measured using C.I.E. standard illuminant "C" with a 2° observer, following the procedures established in ASTM E 308-90. The total solar ultraviolet transmittance (TSUV) is measured over the wavelength range of 300 to 400 nanometers, total solar infrared transmittance (TSIR) is measured over the wavelength range of 720 to 2000 nanometers, and total solar energy
25 transmittance (TSET) is measured over the wavelength range of 300 to 2000 nanometers. The TSUV, TSIR and TSET transmittance data is calculated using Parry Moon air mass 2.0 direct solar irradiance data and integrated using the Trapezoidal Rule, as is known in the art. The shading coefficient

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- (SC), which is the ratio of the total solar heat gain for a sample to the total solar heat gain of a nominal 3 mm thick (1/8 inch) clear glass reference, is calculated using the Window 4.1 computer program available from Lawrence Berkeley Laboratory. The spectral properties presented in Tables 4-6 for
- 5 Examples 17-29 of Table 2 are based on the same wavelength ranges and calculation procedures.

Sample Preparation

The information provided in Table 1 is based on experimental laboratory melts having approximately the following basic batch components:

10	sand	500 gm
	soda ash	162.8 gm
	limestone	42 gm
	dolomite	121 gm
	salt cake	2.5 gm
15	rouge	as required
	Co ₃ O ₄	as required

- Coal was added to each melt to control redox. In preparing the melts, the ingredients were weighed out and blended in a mixer. Half of the material was then placed in a refractory silica crucible and heated to 2650°F (1454°C)
- 20 for 30 minutes. The remaining material was then added to the crucible and heated to 2650°F (1454°C) for 1.5 hours. Next, the molten glass was fritted in water, dried and reheated at 2650°F (1454°C) for one hour. The molten glass was then fritted a second time in water, dried and reheated to 2650°F (1454°C) for two hours. The molten glass was then poured out of the crucible
- 25 and annealed. Samples were cut from the slab and ground and polished for analysis.

The chemical analysis of the glass compositions was determined using a RIGAKU 3370 X-ray fluorescence spectrophotometer. The FeO content for

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Examples 1-16 was determined using wet chemistry techniques, as are well known in the art. The FeO content for Examples 30-34 was calculated using the glass color and spectral performance computer model discussed earlier. The spectral characteristics of the glass were determined on annealed
 5 samples using a Perkin-Elmer Lambda 9 UV/VIS/NIR spectrophotometer prior to tempering the glass or prolonged exposure to ultraviolet radiation, which will affect the spectral properties as will be discussed later.

The following is representative of the basic oxides of the glasses disclosed in Tables 1 and 3:

	<u>Ex. 1-16 of Table 1</u>	<u>Ex. 30-34 of Table 3</u>
SiO ₂	73.3 to 73.9 wt. %	~72.5 wt. %
Na ₂ O	13.2 to 13.6 wt. %	~13.8 wt. %
K ₂ O	0.031 to 0.034 wt. %	~0.046 wt. %
CaO	8.5 to 8.8 wt. %	~9 wt. %
MgO	3.6 to 3.8 wt. %	~3.2 wt. %
Al ₂ O ₃	0.12 to 0.16 wt. %	~0.24 wt. %

10

It is expected that the basic oxide constituents of commercial soda-lime-silica glass compositions based on the experimental melts disclosed in Table 1 and the modeled compositions disclosed in Table 2 would be similar to those discussed earlier.

15

Although not included in Table 1, analysis of Examples 1-16 indicated up to about 6 PPM Cr₂O₃, up to about 30 PPM MnO₂ and up to about 0.013 weight percent TiO₂. These materials were most likely introduced into the glass composition from the batch raw materials or glass processing equipment. Similarly, although not included in Table 3, analysis of Examples
 20 30-34 indicated up to about 9 PPM Cr₂O₃, up to about 26 PPM MnO₂ and up to about 0.021 weight percent TiO₂. These materials were also most likely

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introduced into the glass composition from the batch raw materials or glass processing equipment as well as from residual materials in the glass melter. All of the computer modeled compositions in Table 2 were modeled to include 38 PPM MnO_2 (not shown in Table 2) and Examples 17-23 were modeled to include 7 PPM Cr_2O_3 , which are typical detectable levels of chromium and manganese. The chromium and manganese at these levels were included in the modeled compositions to provide a more accurate representation of the glass using the model. It is believed that these amounts of chromium, manganese and titanium in Examples 1-29 of Tables 1 and 2 and the chromium and manganese in Examples 30-34 of Table 3 are tramp and/or residual levels which will not materially affect the color and spectral properties of the glass. The TiO_2 level in Examples 30-34, which is believed to be due to residual titanium still in the glass melter, should have only a minimal, if any, affect on the glass color and spectral properties of the glass compositions of the instant invention. Furthermore, it is believed that glass compositions embodying the features of the instant invention may be produced with no or trace levels of TiO_2 .

TABLE 1

Experimental Glass Melt Compositions

	Ex. 1	Ex. 2	Ex. 3	Ex. 4	Ex. 5	Ex. 6	Ex. 7	Ex. 8	Ex. 9	Ex. 10	Ex. 11	Ex. 12	Ex. 13	Ex. 14	Ex. 15	Ex. 16
Total Iron (wt.%)	0.591	0.607	0.613	0.616	0.629	0.632	0.637	0.641	0.659	0.661	0.684	0.671	0.679	0.699	0.717	0.720
Fe ₂ O ₃ (wt.%)	0.271	0.311	0.306	0.360	0.371	0.279	0.368	0.337	0.310	0.334	0.333	0.286	0.350	0.399	0.410	0.340
FeO (wt.%)	0.288	0.266	0.276	0.230	0.232	0.318	0.242	0.274	0.314	0.294	0.298	0.258	0.296	0.270	0.276	0.342
Redox	0.487	0.438	0.450	0.373	0.369	0.503	0.380	0.427	0.476	0.445	0.449	0.385	0.436	0.386	0.385	0.475
CoO (PPM)	13	13	7	12	6	14	6	12	7	6	14	12	6	6	6	6

TABLE 2

Computer Modeled Glass Compositions

	Ex. 17	Ex. 18	Ex. 19	Ex. 20	Ex. 21	Ex. 22	Ex. 23	Ex. 24	Ex. 25	Ex. 26	Ex. 27	Ex. 28	Ex. 29
Total Iron (wt.%)	0.45	0.50	0.55	0.75	0.80	0.90	1.0	0.50	0.50	0.50	0.55	0.55	0.55
Fe ₂ O ₃ (wt.%)	0.141	0.211	0.269	0.450	0.480	0.540	0.600	0.211	0.211	0.211	0.269	0.269	0.269
FeO (wt.%)	0.278	0.260	0.253	0.270	0.288	0.324	0.360	0.260	0.260	0.260	0.253	0.253	0.253
Redox	0.58	0.52	0.46	0.36	0.36	0.36	0.36	0.52	0.52	0.52	0.46	0.46	0.46
CoO (PPM)	5	5	5	5	3	3	3	5	5	5	5	5	5
Cr ₂ O ₃ (PPM)	7	7	7	7	7	7	7	25	50	75	25	50	75

TABLE 3
Production Glass Compositions

	Ex. 30	Ex. 31	Ex. 32	Ex. 33	Ex. 34
Total Iron (wt. %)	0.55	0.554	0.552	0.553	0.548
Fe ₂ O ₃ (wt. %)	0.263	0.258	0.254	0.256	0.252
FeO (wt. %)	0.258	0.266	0.268	0.267	0.266
Redox	0.469	0.480	0.485	0.482	0.486
CoO (PPM)	11	9	9	9	10

TABLE 4
Spectral Properties at 0.084 inches (2.13 mm)

	Ex. 1	Ex. 2	Ex. 3	Ex. 4	Ex. 5	Ex. 6	Ex. 7	Ex. 8	Ex. 9	Ex. 10	Ex. 11	Ex. 12	Ex. 13	Ex. 14	Ex. 15	Ex. 16	Ex. 17
LTA (%)	77.8	78.3	78.7	79.4	80.4	76.5	79.7	78.3	77.8	77.5	76.9	78.4	78.0	78.7	78.4	76.0	80.4
LTS (%)	79.3	79.7	80.2	80.6	81.6	78.1	81.0	79.7	79.3	79.12	78.4	79.7	79.5	80.1	79.8	77.7	81.7
TSUV (%)	69.0	68.3	68.2	66.4	65.3	68.2	65.8	66.5	67.2	67.0	66.9	65.0	65.9	64.2	63.4	65.2	75.1
TSIR (%)	32.9	34.5	33.2	39.4	41.1	29.0	38.0	35.5	30.4	29.8	29.7	35.8	31.8	35.1	34.4	27.0	39.4
TSET (%)	54.0	55.0	54.4	58.1	59.2	51.3	57.3	55.6	52.5	52.0	51.9	55.7	53.3	55.3	54.7	49.8	58.8
DW (nm)	487.1	487.2	487.7	487.7	488.8	487.3	488.4	487.5	487.8	487.8	487.2	487.9	488.0	488.6	488.7	487.9	486.4
Pe (%)	6.30	5.98	5.86	4.99	4.33	6.78	4.85	5.64	6.18	6.31	6.51	5.30	5.85	5.15	5.13	6.67	5.28
SC	0.74	0.75	0.75	0.78	0.79	0.72	0.77	0.76	0.73	0.72	0.72	0.76	0.74	0.75	0.75	0.71	0.78

	Ex. 18	Ex. 19	Ex. 20	Ex. 21	Ex. 22	Ex. 23	Ex. 24	Ex. 25	Ex. 26	Ex. 27	Ex. 28	Ex. 29	Ex. 30	Ex. 31	Ex. 32	Ex. 33	Ex. 34
LTA (%)	80.3	80.4	79.3	78.9	77.6	76.3	80.1	79.9	79.6	80.2	79.9	79.7	79.8	79.7	79.7	79.8	79.9
LTS (%)	81.6	81.6	80.5	80.2	78.9	77.7	81.4	81.1	80.9	81.4	81.1	80.9	81.0	81.2	81.0	81.0	81.1
TSUV (%)	72.2	69.3	61.8	60.8	58.7	56.8	72.2	72.2	72.1	69.3	69.3	69.2	66.0	66.0	66.3	66.4	66.9
TSIR (%)	39.6	40.4	38.3	36.2	32.4	29.0	39.5	39.5	39.5	40.4	40.4	40.4	39.8	38.8	38.6	38.8	38.9
TSET (%)	58.7	59.1	57.1	55.7	53.0	50.5	58.6	58.4	58.2	58.9	58.8	58.6	58.5	57.9	57.8	57.9	58.0
DW (nm)	487.0	487.8	489.9	490.3	490.2	490.1	487.4	487.9	488.4	488.2	488.8	489.4	488.2	488.1	488.1	488.0	488.0
Pe (%)	5.02	4.59	4.14	4.25	4.78	5.31	4.96	4.88	4.80	4.53	4.45	4.38	4.63	4.68	4.70	4.75	4.77
SC	0.78	0.78	0.77	0.76	0.74	0.71	0.78	0.78	0.78	0.78	0.78	0.78	0.78	0.77	0.77	0.77	0.78

TABLE 5

Spectral Properties at 0.154 inches (3.9 mm)

	Ex. 1	Ex. 2	Ex. 3	Ex. 4	Ex. 5	Ex. 6	Ex. 7	Ex. 8	Ex. 9	Ex. 10	Ex. 11	Ex. 12	Ex. 13	Ex. 14	Ex. 15	Ex. 16	Ex. 17
LTA (%)	68.3	69.0	69.7	70.7	72.3	66.2	71.3	69.1	68.2	67.9	66.8	69.1	68.7	69.7	69.2	65.5	72.4
LTS (%)	70.7	71.3	72.1	72.8	74.3	68.8	73.4	71.3	70.7	70.4	69.3	71.3	71.0	71.8	71.4	68.1	74.5
TSUV (%)	59.2	58.3	58.4	55.9	54.6	58.0	55.3	55.9	56.9	56.7	56.4	53.9	55.2	53.1	52.0	54.2	67.6
TSIR (%)	14.9	16.2	15.1	20.3	21.8	12.1	19.1	17.0	13.0	12.6	12.8	17.3	14.0	16.7	16.0	10.7	20.3
TSET (%)	40.0	40.9	40.5	43.8	45.0	37.4	43.1	41.2	38.6	38.2	38.1	41.4	39.3	41.0	40.4	36.1	44.9
DW (nm)	486.8	486.9	487.5	487.5	488.6	487.0	488.2	487.3	487.6	487.5	486.9	487.7	487.8	488.4	488.6	487.6	486.1
Pe (%)	11.22	10.69	10.44	8.96	7.79	12.04	8.71	10.09	10.98	11.20	11.58	9.48	10.42	9.21	9.17	11.81	9.49
SC	0.62	0.63	0.63	0.66	0.67	0.60	0.65	0.63	0.61	0.61	0.61	0.64	0.62	0.63	0.63	0.59	0.66

	Ex. 18	Ex. 19	Ex. 20	Ex. 21	Ex. 22	Ex. 23	Ex. 24	Ex. 25	Ex. 26	Ex. 27	Ex. 28	Ex. 29	Ex. 30	Ex. 31	Ex. 32	Ex. 33	Ex. 34
LTA (%)	72.2	72.4	70.6	70.0	67.9	65.8	71.9	71.5	71.1	72.1	71.6	71.2	71.5	71.2	71.3	71.4	71.5
LTS (%)	74.3	74.3	72.5	72.0	70.0	68.1	74.0	73.6	73.1	74.0	73.6	73.2	73.4	73.2	73.3	73.4	73.5
TSUV (%)	63.7	59.8	50.1	48.7	46.2	43.8	63.6	63.6	63.5	59.8	59.7	59.6	55.8	55.7	56.0	56.2	56.8
TSIR (%)	20.4	21.2	19.3	17.5	14.4	11.9	20.4	20.4	20.3	21.2	21.2	21.1	20.7	19.8	19.6	19.7	19.8
TSET (%)	44.7	44.9	42.6	41.2	38.5	36.2	44.5	44.2	43.9	44.7	44.5	44.2	44.3	43.7	43.6	43.7	43.8
DW (nm)	486.8	487.6	489.8	490.1	490.1	490.0	487.1	487.7	488.3	488.0	488.6	489.2	488.0	487.9	487.9	487.8	487.8
Pe (%)	9.02	8.25	7.43	7.63	8.54	9.43	8.90	8.75	8.59	8.15	7.99	7.85	8.14	8.41	8.45	8.53	8.55
SC	0.66	0.66	0.64	0.63	0.61	0.59	0.66	0.66	0.66	0.66	0.66	0.66	0.66	0.65	0.65	0.65	0.65

TABLE 6
Spectral Properties at 0.223 inches (5.5 mm)

	Ex. 1	Ex. 2	Ex. 3	Ex. 4	Ex. 5	Ex. 6	Ex. 7	Ex. 8	Ex. 9	Ex. 10	Ex. 11	Ex. 12	Ex. 13	Ex. 14	Ex. 15	Ex. 16	Ex. 17
LTA (%)	60.1	61.0	61.9	63.2	65.2	57.6	63.9	61.1	60.1	59.7	58.3	61.2	60.6	61.9	61.3	56.7	65.4
LTS (%)	63.2	63.9	65.0	65.8	67.7	60.8	66.6	64.0	63.2	62.8	61.4	63.9	63.6	64.6	64.1	59.9	68.1
TSUV (%)	51.8	50.9	50.9	48.1	46.6	50.3	47.4	48.1	49.2	49.0	48.5	45.7	47.3	44.9	43.6	46.0	61.6
TSIR (%)	7.12	7.93	7.21	10.9	11.9	5.4	9.9	8.5	5.9	5.6	5.9	8.8	6.5	8.3	7.8	4.5	10.8
TSET (%)	32.0	32.8	32.6	35.1	36.3	29.9	34.7	33.0	31.0	30.7	30.4	33.0	31.5	32.8	32.2	28.7	36.6
DW (nm)	486.6	486.7	487.3	487.3	488.5	486.8	488.1	487.1	487.4	487.3	486.7	487.5	487.6	488.2	488.4	487.4	485.9
Pe (%)	15.82	15.09	14.72	12.72	11.07	16.90	12.34	14.26	15.45	15.75	16.29	13.42	14.68	13.02	12.96	16.55	13.48
SC	0.55	0.56	0.56	0.58	0.59	0.54	0.58	0.56	0.54	0.54	0.54	0.56	0.55	0.56	0.56	0.53	0.59

	Ex. 18	Ex. 19	Ex. 20	Ex. 21	Ex. 22	Ex. 23	Ex. 24	Ex. 25	Ex. 26	Ex. 27	Ex. 28	Ex. 29	Ex. 30	Ex. 31	Ex. 32	Ex. 33	Ex. 34
LTA (%)	65.1	65.3	63.0	62.3	59.6	57.1	64.8	64.2	63.7	64.9	64.3	63.8	64.1	63.8	63.9	64.0	64.2
LTS (%)	67.8	67.8	65.5	64.8	62.3	59.9	67.4	66.8	66.3	67.4	66.9	66.3	66.6	66.3	66.4	66.6	66.8
TSUV (%)	57.0	52.5	41.5	40.0	37.2	34.7	56.9	56.8	56.7	52.4	52.3	52.2	48.2	48.0	48.4	48.6	49.3
TSIR (%)	10.8	1.4	10.0	8.8	6.7	5.2	10.8	10.8	10.8	11.4	11.4	11.4	11.1	10.4	10.3	10.4	10.4
TSET (%)	36.3	36.4	33.9	32.7	30.4	28.3	36.0	35.6	35.3	36.1	35.7	35.4	35.6	35.1	35.1	35.2	35.3
DW (nm)	486.5	487.4	489.7	490.1	490.0	489.9	486.9	487.5	488.1	487.8	488.5	489.1	487.9	487.7	487.7	487.7	487.6
Pe (%)	12.80	11.73	10.53	10.80	12.04	13.24	12.64	12.39	12.16	11.56	11.33	11.11	11.57	11.93	12.0	12.11	12.14
SC	0.59	0.59	0.57	0.56	0.54	0.52	0.59	0.58	0.58	0.59	0.59	0.58	0.58	0.58	0.58	0.58	0.58

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Referring to Tables 1-5, the present invention provides a blue colored glass using a standard soda-lime-silica glass base composition and additionally iron and cobalt, and optionally chromium, as infrared and ultraviolet radiation absorbing materials and colorants. In particular, the

5 infrared and ultraviolet radiation absorbing and colorant portion of the glass includes about 0.40 to 1.0 wt.% total iron, preferably about 0.50 to 0.75 wt.% total iron, and about 4 to 40 PPM CoO, preferably about 4 to 20 PPM CoO. In addition, the composition may include up to 100 PPM Cr₂O₃, preferably 25 to 50 PPM. The glass has a redox of greater than 0.35 up to 0.60, preferably

10 from about 0.36 to 0.50. In one embodiment of the invention, the glass has a luminous transmittance (LTA) of at least 55%, a dominant wavelength of between about 485 to 489 nm, preferably 487 to 489 nm. In another embodiment of the invention, glass has a luminous transmittance of at least 65% at a thickness of 0.154 inches and a dominant wavelength of about 485

15 to 492 nm, preferably about 487 to 489 nm. The glass has an excitation purity of about 3 to 18%. However, it should be appreciated that excitation purity depends on the glass thickness. As a result, it is preferred that glass of the present invention having a thickness of about 0.071 to 0.126 inches (1.8 to 3.2 mm) have a Pe of about 3 to 8%; a thickness of about 0.126 to 0.189

20 inches (3.2 to 4.9 mm) have a Pe of about 5 to 12%; and a thickness of about 0.189 to 0.315 inches (4.9 to 8 mm) have a Pe of about 10 to 18%.

In another embodiment of the instant, the infrared and ultraviolet radiation absorbing materials and colorants of the glass include about 0.50 to 0.60 wt.% total, about 4 to 12 PPM CoO, 0 to about 100 PPM Cr₂O₃, and 0 to

25 about 0.50 wt.% TiO₂, and a redox of about 0.45 to 0.50. In one particular embodiment, the glass includes only trace amounts of Cr₂O₃ and up to about 0.021 wt.% TiO₂. In addition, the glass has a luminous transmittance (LTA) of at least 70%, a TSUV of no greater than 60%, a TSIR of no greater than

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about 30%, and/or a TSET of no greater than 50% at a thickness of about 0.154 inches. The color of the glass is characterized by a dominant wavelength of between about 487 to 489 nm and an excitation purity of about 7-10%.

5 Glass compositions as disclosed herein and made by the float process typically range from a sheet thickness of about 1 millimeter to 10 millimeters.

For the vehicle glazing applications, it is preferred that the glass sheets having a composition as disclosed herein have a thickness within the range of 0.071 to 0.197 inches (1.8 to 5 mm). It is anticipated that when using a single
10 glass ply, the glass will be tempered, e.g. for an automotive side or rear window, and when multiple plies are used, the glass will be annealed and laminated together using a thermoplastic adhesive, e.g. an automobile windshield which laminates two annealed glass plies together using a polyvinyl butyral interlayer, wherein at least one of the glass plies is a glass
15 sheet having a composition as disclosed herein. In addition, when the glass is used in selected areas of a motor vehicle, e.g. the windshield and front door windows and in some instances the rear window, it is required that the glass have an LTA of at least 70%. In addition, the glass compositions disclosed in the present invention should have a TSUV of no greater than
20 60%, preferably no greater than 57%, a TSIR of no greater than about 35%, preferably no greater than about 30%, and/or a TSET of no greater than about 55%, preferably, no greater than about 50% at a thickness of about 0.154 inches.

In architectural glazing applications, there is generally no legal
25 requirement with respect to solar transmittance (LTS) of the glass; however, in the present invention, it is preferred that the glass have an LTS of about 60 to 70%, preferably about 63 to 67% and a shading coefficient of no greater

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than about 0.70, and preferably no greater than about 0.65 at a thickness of about 0.223 inches.

It is expected that the spectral properties of the glass will change after tempering the glass and further upon prolonged exposure to ultraviolet radiation, commonly referred to as solarization. In particular, it is estimated that tempering and solarization of the glass compositions disclosed herein will reduce LTA, LTS and TSIR by about 0.5 to 1%, reduce the TSUV by about 1 to 2%, and TSET by about 1 to 1.5%. As a result, in one embodiment of the invention, the glass composition has selected spectral properties that initially fall outside the desired ranges previously discussed but fall within the desired ranges after tempering and/or solarization.

It should also be appreciated that since the luminous and solar transmittance (LTA and LTS) is reduced by these conditions, in order to maintain the transmittance above a desired minimum level, the initial LTA or LTS of the glass after production should be sufficiently high so that any losses attributable to tempering and solarization will not reduce the transmittance to an unacceptable level.

Vanadium may be used as a partial or complete replacement for the chromium in the glass compositions of the present invention. More specifically, vanadium, which is expressed herein in terms of V_2O_5 , imparts a yellow-green color to the glass and absorbs both ultraviolet and infrared radiation at different valence states. It is believed that the 100 PPM Cr_2O_3 as discussed above may be completely replaced by about 400 PPM V_2O_5 .

As discussed earlier, other materials may also be added to the glass compositions disclosed herein to further reduce infrared and ultraviolet radiation transmission and/or control glass color. Inclusion of chromium and manganese has been discussed earlier. It is contemplated that the following

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materials may also be added to the iron and cobalt containing soda-lime-silica glass disclosed herein:

	SnO ₂	0 to about 2.0 wt. %
	CeO ₂	0 to about 1.0 wt. %
5	TiO ₂	0 to about 0.5 wt. %
	ZnO	0 to about 0.5 wt. %
	Nd ₂ O ₃	0 to about 0.5 wt. %
	MoO ₃	0 to about 100 PPM
	NiO	0 to about 10 PPM
10	Se	0 to about 3 PPM

As should be appreciated, adjustment may have to be made to the basic constituents of the glass to account for any coloring and/or redox affecting power of these additional materials.

Other variations as are known to those of skill in the art may be
15 resorted to without departing from the scope of the invention as defined by the claims that follow.

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WE CLAIM:

1. A blue colored, infrared and ultraviolet radiation absorbing glass composition having a base glass portion comprising:

5	SiO ₂	about 66 to 75 percent by weight
	Na ₂ O	about 10 to 20 percent by weight
	CaO	about 5 to 15 percent by weight
	MgO	0 to about 5 percent by weight
	Al ₂ O ₃	0 to about 5 percent by weight
10	K ₂ O	0 to about 5 percent by weight

and a solar radiation absorbing and colorant portion consisting essentially of:

total iron	about 0.40 to 1.0 percent by weight
CoO	about 4 to 40 PPM
Cr ₂ O ₃	0 to about 100 PPM,

15 the glass having a redox of greater than 0.35 up to about 0.60, a luminous transmittance of at least 55 percent, and a color characterized by a dominant wavelength of 485 to 489 nanometers and an excitation purity of about 3 to 18 percent.

20 2. The composition as in claim 1 wherein the total iron concentration is from about 0.50 to 0.75 weight percent and the redox is from about 0.36 to 0.50.

25 3. The composition as in claim 2 wherein the CoO concentration is from about 4 to 20 PPM.

4. The composition as in claim 3 wherein the color of the glass is characterized by a dominant wavelength in the range of about 487 to 489

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nanometers and an excitation purity of about 3 to 8 percent at at least one thickness ranging from about 0.071 to 0.126 inches, an excitation purity of about 5 to 12 percent at at least one thickness ranging from about 0.126 to 0.189 inches, and an excitation purity of about 10 to 18 percent at at least one thickness ranging from about 0.189 to 0.315 inches.

5 5. The composition as in claim 4 wherein the glass has a luminous transmittance of about 70 percent or greater.

10 6. The composition as in claim 4 wherein the glass has a solar transmittance of about 55 to 70 percent and a shading coefficient of about 0.70 or less.

 7. The composition as in claim 1 wherein the CoO concentration is
15 from about 4 to 20 PPM.

 8. The composition as in claim 1 wherein the Cr₂O₃ concentration is from about 25 to 50 PPM.

20 9. The composition as in claim 1 wherein the glass has a luminous transmittance of about 70 percent or greater.

 10. The composition as in claim 9 wherein the glass has a total solar ultraviolet transmittance of about 60 percent or less, a total solar
25 infrared transmittance of about 35 percent or less and a total solar energy transmittance of about 55 percent or less at a thickness of about 0.154 inches.

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11. The composition as in claim 10 wherein the glass has a total solar ultraviolet transmittance of about 57 percent or less, a total solar infrared transmittance of about 30 percent or less and a total solar energy transmittance of about 50 percent or less at a thickness of about 0.154 inches.

12. The composition as in claim 1 wherein the glass has a solar transmittance of about 55 to 70 percent and a shading coefficient of about 0.70 or less at a thickness of about 0.223 inches.

13. The composition as in claim 12 wherein the glass has a solar transmittance of about 63 to 67 percent and a shading coefficient of about 0.65 or less at a thickness of about 0.223 inches.

14. The composition as in claim 1 wherein the color of the glass is characterized by a dominant wavelength in the range of about 487 to 489 nanometers.

15. The composition as in claim 1 wherein the color of the glass is characterized by an excitation purity of about 3 to 8 percent at at least one thickness ranging from about 0.071 to 0.126 inches, an excitation purity of about 5 to 12 percent at at least one thickness ranging from about 0.126 to 0.189 inches, and an excitation purity of about 10 to 18 percent at at least one thickness ranging from about 0.189 to 0.315 inches.

16. A glass sheet formed by the float process from the glass composition as recited in claim 1.

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17. An automobile windshield made from at least one glass sheet of claim 16.

18. The glass sheet as in claim 16 wherein the sheet has a thickness between 1.8 to 10 mm.

19. A blue colored, infrared and ultraviolet radiation absorbing glass composition having a base glass portion comprising:

10	SiO ₂	about 66 to 75 percent by weight
	Na ₂ O	about 10 to 20 percent by weight
	CaO	about 5 to 15 percent by weight
	MgO	0 to about 5 percent by weight
	Al ₂ O ₃	0 to about 5 percent by weight
15	K ₂ O	0 to about 5 percent by weight
	and a solar radiation absorbing and colorant portion consisting essentially of:	
	total iron	about 0.40 to 1.0 percent by weight
	CoO	about 4 to 40 PPM
	SnO ₂	0 to about 2.0 percent by weight
20	CeO ₂	0 to about 1.0 percent by weight
	TiO ₂	0 to about 0.5 percent by weight
	ZnO	0 to about 0.5 percent by weight
	Nd ₂ O ₃	0 to about 0.5 percent by weight
	MnO ₂	0 to about 0.1 percent by weight
25	MoO ₃	0 to about 100 PPM
	Cr ₂ O ₃	0 to about 100 PPM,
	V ₂ O ₅	0 to about 400 PPM
	NiO	0 to about 10 PPM
	Se	0 to about 3 PPM

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the glass having a redox greater than 0.35 up to about 0.60, a luminous transmittance of at least 55 percent, and a color characterized by a dominant wavelength of 485 to 489 nanometers and an excitation purity of about 3 to 18 percent.

5

20. The composition as in claim 19 wherein the color of the glass is characterized by a dominant wavelength in the range of about 487 to 489 nanometers and an excitation purity of about 3 to 8 percent at at least one thickness in the range from about 0.071 to 0.126 inches, an excitation purity of about 5 to 12 percent at at least one thickness in the range from about 0.126 to 0.189 inches, and an excitation purity of about 10 to 18 percent at at least one thickness in the range from about 0.189 to 0.315 inches.

10

21. A blue colored, infrared and ultraviolet radiation absorbing glass composition having a base glass portion comprising:

15

SiO ₂	about 66 to 75 percent by weight
Na ₂ O	about 10 to 20 percent by weight
CaO	about 5 to 15 percent by weight
MgO	0 to about 5 percent by weight
Al ₂ O ₃	0 to about 5 percent by weight
K ₂ O	0 to about 5 percent by weight

20

and a solar radiation absorbing and colorant portion consisting essentially of:

total iron	about 0.40 to 1.0 percent by weight
CoO	about 4 to 40 PPM
Cr ₂ O ₃	0 to about 100 PPM,

25

the glass having a redox of greater than 0.35 up to about 0.60, a luminous transmittance of at least 65% at a thickness of about 0.154 inches, and a

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color characterized by a dominant wavelength of about 485 to 492 nanometers and an excitation purity of about 3 to 18 percent.

22. The composition as in claim 21 wherein the total iron
5 concentration is from about 0.50 to 0.75 weight percent, the CoO concentration is from about 4 to 20 PPM, and the redox is from about 0.36 to 0.50, and the color of the glass is characterized by a dominant wavelength in the range of about 487 to 489 nanometers and an excitation purity of about 3 to 8 percent at at least one thickness ranging from about 0.071 to 0.126
10 inches, an excitation purity of about 5 to 12 percent at at least one thickness ranging from about 0.126 to 0.189 inches, and an excitation purity of about 10 to 18 percent at at least one thickness ranging from about 0.189 to 0.315 inches.

15 23. The composition as in claim 22 wherein the glass has a total solar ultraviolet transmittance of about 60 percent or less, a total solar infrared transmittance of about 35 percent or less and a total solar energy transmittance of about 55 percent or less at a thickness of about 0.154 inches.

20

24. A blue colored, infrared and ultraviolet radiation absorbing glass composition having a base glass portion comprising:

25	SiO ₂	about 66 to 75 percent by weight
	Na ₂ O	about 10 to 20 percent by weight
	CaO	about 5 to 15 percent by weight
	MgO	0 to about 5 percent by weight
	Al ₂ O ₃	0 to about 5 percent by weight
	K ₂ O	0 to about 5 percent by weight

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and a solar radiation absorbing and colorant portion consisting essentially of:

	total iron	about 0.40 to 1.0 percent by weight
	CoO	about 4 to 40 PPM
5	SnO ₂	0 to about 2.0 percent by weight
	CeO ₂	0 to about 1.0 percent by weight
	TiO ₂	0 to about 0.5 percent by weight
	ZnO	0 to about 0.5 percent by weight
	Nd ₂ O ₃	0 to about 0.5 percent by weight
	MnO ₂	0 to about 0.1 percent by weight
10	MoO ₃	0 to about 100 PPM
	Cr ₂ O ₃	0 to about 100 PPM,
	V ₂ O ₅	0 to about 400 PPM
	NiO	0 to about 10 PPM
	Se	0 to about 3 PPM

- 15 the glass having a redox of greater than 0.35 up to about 0.60, a luminous transmittance of at least 65% at a thickness of about 0.154 inches, and a color characterized by a dominant wavelength of about 485 to 492 nanometers and an excitation purity of about 3 to 18 percent.

- 20 25. The composition as in claim 24 wherein the color of the glass is characterized by a dominant wavelength in the range of about 487 to 489 nanometers and an excitation purity of about 3 to 8 percent at at least one thickness ranging from about 0.071 to 0.126 inches, an excitation purity of about 5 to 12 percent at at least one thickness ranging from about 0.126 to
- 25 0.189 inches, and an excitation purity of about 10 to 18 percent at at least one thickness ranging from about 0.189 to 0.315 inches.

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26. A blue colored, infrared and ultraviolet radiation absorbing glass composition having a base glass portion comprising:

5	SiO_2	about 66 to 75 percent by weight
	Na_2O	about 10 to 20 percent by weight
	CaO	about 5 to 15 percent by weight
	MgO	0 to about 5 percent by weight
	Al_2O_3	0 to about 5 percent by weight
	K_2O	0 to about 5 percent by weight

and a solar radiation absorbing and colorant portion consisting essentially of:

10	total iron	about 0.50 to 0.60 percent by weight
	CoO	about 4 to 12 PPM
	Cr_2O_3	0 to about 100 PPM
	TiO_2	0 to about 0.50 percent by weight,

the glass having a redox of about 0.45 to 0.50, a luminous transmittance of at least 70 % at a thickness of about 0.154 inches, and a color characterized by a dominant wavelength of about 487 to 489 nanometers and an excitation purity of about 7 to 10 percent.

27. The composition as in claim 26 wherein the glass has a total solar ultraviolet transmittance of about 60 percent or less, a total solar infrared transmittance of about 30 percent or less and a total solar energy transmittance of about 50 percent or less at a thickness of about 0.154 inches.

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 98/21987

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C03C3/087 C03C4/02 C03C4/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C03C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	EP 0 820 964 A (PILKINGTON PLC) 28 January 1998 see page 3, line 11 - page 5, line 4; examples 3,6,14,17,20 ----	1-7,9-27
X	US 2 755 212 A (BROWN) 17 July 1956 see the whole document ----	1-7,9-27
A	US 4 792 536 A (PECORARO GEORGE A ET AL) 20 December 1988 cited in the application see claims; examples; tables ----	1-27
A	US 5 344 798 A (MORIMOTO SHIGEKI ET AL) 6 September 1994 cited in the application see claims; examples ----- -/--	1-27

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

3 February 1999

Date of mailing of the international search report

11/02/1999

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 98/21987

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, A	EP 0 814 064 A (PPG INDUSTRIES INC) 29 December 1997 see page 2, line 53 - page 4, line 18 -----	1-27

INTERNATIONAL SEARCH REPORT

Information on patent family members

Interr. Application No

PCT/US 98/21987

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0820964 A	28-01-1998	JP 10101367 A	21-04-1998
US 2755212 A	17-07-1956	NONE	
US 4792536 A	20-12-1988	AU 599494 B	19-07-1990
		AU 1852088 A	05-01-1989
		AU 622327 B	02-04-1992
		AU 5593690 A	13-09-1990
		CA 1294990 A	28-01-1992
		EP 0297404 A	04-01-1989
		GR 3001585 T	23-11-1992
		JP 1018938 A	23-01-1989
		JP 5027578 B	21-04-1993
		JP 7165435 A	27-06-1995
		MX 9205146 A	31-03-1994
US 5344798 A	06-09-1994	JP 2544035 B	16-10-1996
		JP 5043266 A	23-02-1993
		DE 69215159 D	19-12-1996
		DE 69215159 T	06-03-1997
		EP 0527487 A	17-02-1993
EP 0814064 A	29-12-1997	US 5688727 A	18-11-1997
		AU 685370 A	15-01-1998
		BR 9702479 A	15-09-1998
		CA 2206826 A	17-12-1997
		CN 1172777 A	11-02-1998
		JP 10067538 A	10-03-1998
		NZ 314790 A	24-09-1998